

Appl. No. 10/528,082  
Amdt. dated May 19, 2008  
Reply to Office Action of January 17, 2008

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**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings of claims in the application:

**Listing of Claims:**

1. (Currently amended) A process ~~for treating a cancer patient~~ to induce in ~~said patient~~ an effector cell mediated immune response against ~~the cancerous tumor cells in a cancer patient~~, said method comprising[[:] ]

administering, to a cancer patient, a ~~tumor-derived~~ biologically generated virus or virus-like particle with a cellular membrane from a host cell, said membrane comprising  
an MHC molecule that presents that have been modified to mimic cells capable of  
presenting one or more tumor specific antigens, and  
a co-stimulatory molecule,

wherein said administering is of to a mammalian immune system in an amount  
effective to induce an effector cell mediated immune response against the ~~cancerous tumor cells~~  
~~in said patient, whereby in the cancer patient the immune response would reduce the amount of~~  
~~cancerous cells.~~

2. (Currently amended) The process of claim 1 wherein said immune  
response is mediated by effector cells are T cells.

3. (Currently amended) The process of claim 1 wherein said ~~tumor-derived~~  
biologically generated particles are released from homologous tumor cells ~~derived~~ from the  
patient.

4. (Currently amended) The process of claim 1 wherein said ~~tumor-derived~~  
biologically generated particles are released from matched major histocompatibility complex  
containing tumor cells.

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5. (Currently amended) The process of claim 1 wherein said ~~tumor-derived~~ biologically generated particles are released from non-homologous tumor cell lines containing one or more matched human leukocyte antigens.

6. (Original) The process of claim 1 wherein said particles are generated as virus-like-particles.

7. (Original) The process of claim 1 wherein said particles are generated as inactivated intact virus particles.

8. (Currently amended) ~~[[The]] A process of claim 1 for treating a cancer patient to induce in said patient an effector cell immune response against the cancerous cells,~~  
comprising; administering to a cancer patient tumor-derived biologically generated particles that have been modified to mimic cells capable of presenting antigens to a mammalian immune system in an amount effective to induce an immune response against the cancerous cells, whereby in the cancer patient the immune response would reduce the amount of cancerous cells, and

wherein said particles mimic dendritic cells.

9-15. (canceled)

16. (New) The process of claim 1 wherein said immune response reduces the number of tumor cells in said patient and thereby treats cancer in said patient.

17. (New) The process of claim 1 wherein said host cell expresses one or more tumor specific antigens on the cell's cell membrane and said particle has a membrane that further comprises the one or more antigens.

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18. (New) The process of claim 1 wherein said particle mimics a dendritic cell.
19. (New) A process to induce an effector cell mediated immune response against tumor cells in a cancer patient, said method comprising  
preparing a biologically generated virus or virus-like particle with a membrane from a host cell, said membrane comprising  
an MHC molecule that presents one or more tumor specific antigens, and  
a co-stimulatory molecule;  
administering said particle to a cancer patient in an amount effective to induce an effector cell mediated immune response against tumor cells in said patient.
20. (New) The process of claim 19 wherein said host cell expresses one or more tumor specific antigens on the cell's cell membrane and said particle has a membrane that further comprises the one or more antigens.
21. (New) The process of claim 1 wherein said host cell is a non-tumor cell.
22. (New) The process of claim 19 wherein said host cell is a non-tumor cell.
23. (New) The process of claim 1 wherein said co-stimulatory molecule is a B7 family molecule.
24. (New) The process of claim 17 wherein said co-stimulatory molecule is a B7 family molecule.
25. (New) The process of claim 19 wherein said co-stimulatory molecule is a B7 family molecule.

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26. (New) The process of claim 20 wherein said co-stimulatory molecule is a B7 family molecule.